Clemente Capasso

List of Publications by Citations

1,363 papers	71,124 citations	126 h-index	197 g-index
1,403 ext. papers	77,848 ext. citations	5.2 avg, IF	8.87 L-index

#	Paper	IF	Citations
1363	Carbonic anhydrases: novel therapeutic applications for inhibitors and activators. *Nature Reviews Drug Discovery*, **2008**, 7, 168-81	64.1	2297
1362	Interfering with pH regulation in tumours as a therapeutic strategy. *Nature Reviews Drug Discovery*, **2011**, 10, 767-77	64.1	1146
1361	Carbonic anhydrase inhibitors. *Medicinal Research Reviews*, **2003**, 23, 146-89	14.4	1062
1360	Multiple binding modes of inhibitors to carbonic anhydrases: how to design specific drugs targeting 15 different isoforms?. *Chemical Reviews*, **2012**, 112, 4421-68	68.1	889
1359	Targeting tumor hypoxia: suppression of breast tumor growth and metastasis by novel carbonic anhydrase IX inhibitors. *Cancer Research*, **2011**, 71, 3364-76	10.1	563
1358	Anticancer and antiviral sulfonamides. *Current Medicinal Chemistry*, **2003**, 10, 925-53	4.3	557
1357	Hypoxia activates the capacity of tumor-associated carbonic anhydrase IX to acidify extracellular pH. *FEBS Letters*, **2004**, 577, 439-45	3.8	556
1356	Carbonic anhydrase inhibitors. *Bioorganic and Medicinal Chemistry Letters*, **2010**, 20, 3467-74	2.9	538
1355	Carbonic anhydrases: current state of the art, therapeutic applications and future prospects. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2004**, 19, 199-229	5.6	532
1354	Structure and function of carbonic anhydrases. *Biochemical Journal*, **2016**, 473, 2023-32	3.8	524
1353	Natural products in drug discovery: advances and opportunities. *Nature Reviews Drug Discovery*, **2021**, 20, 200-216	64.1	522
1352	How many carbonic anhydrase inhibition mechanisms exist?. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2016**, 31, 345-60	5.6	485
1351	Structure-based drug discovery of carbonic anhydrase inhibitors. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2012**, 27, 759-72	5.6	483
1350	Carbonic anhydrases as targets for medicinal chemistry. *Bioorganic and Medicinal Chemistry*, **2007**, 15, 4336-50	3.4	462
1349	Non-zinc mediated inhibition of carbonic anhydrases: coumarins are a new class of suicide inhibitors. *Journal of the American Chemical Society*, **2009**, 131, 3057-62	16.4	400
1348	Crystal structure of the catalytic domain of the tumor-associated human carbonic anhydrase IX. *Proceedings of the National Academy of Sciences of the United States of America*, **2009**, 106, 16233-8	11.5	399
1347	Carbonic anhydrases--an overview. *Current Pharmaceutical Design*, **2008**, 14, 603-14	3.3	397

1346	Ureido-substituted benzenesulfonamides potently inhibit carbonic anhydrase IX and show antimetastatic activity in a model of breast cancer metastasis. *Journal of Medicinal Chemistry*, **2011**, 54, 1896-902	8.3	391
1345	Unexpected nanomolar inhibition of carbonic anhydrase by COX-2-selective celecoxib: new pharmacological opportunities due to related binding site recognition. *Journal of Medicinal Chemistry*, **2004**, 47, 550-7	8.3	381
1344	Recent developments in targeting carbonic anhydrase IX for cancer therapeutics. *Oncotarget*, **2012**, 3, 84-97	3.3	325
1343	Targeting tumor-associated carbonic anhydrase IX in cancer therapy. *Trends in Pharmacological Sciences*, **2006**, 27, 566-73	13.2	321
1342	Protease inhibitors of the sulfonamide type: anticancer, antiinflammatory, and antiviral agents. *Medicinal Research Reviews*, **2003**, 23, 535-58	14.4	320
1341	Deciphering the mechanism of carbonic anhydrase inhibition with coumarins and thiocoumarins. *Journal of Medicinal Chemistry*, **2010**, 53, 335-44	8.3	311
1340	Advances in structure-based drug discovery of carbonic anhydrase inhibitors. *Expert Opinion on Drug Discovery*, **2017**, 12, 61-88	6.2	298
1339	An overview of the alpha-, beta- and gamma-carbonic anhydrases from Bacteria: can bacterial carbonic anhydrases shed new light on evolution of bacteria?. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2015**, 30, 325-32	5.6	279
1338	Discovery of a new family of carbonic anhydrases in the malaria pathogen Plasmodium falciparum--the ζ-carbonic anhydrases. *Bioorganic and Medicinal Chemistry Letters*, **2014**, 24, 4389-4396	2.9	258
1337	Carbonic anhydrase inhibitors. Synthesis of water-soluble, topically effective, intraocular pressure-lowering aromatic/heterocyclic sulfonamides containing cationic or anionic moieties: is the tail more important than the ring?. *Journal of Medicinal Chemistry*, **1999**, 42, 2641-50	8.3	250
1336	Carbonic anhydrase activators: X-ray crystallographic and spectroscopic investigations for the interaction of isozymes I and II with histamine. *Biochemistry*, **1997**, 36, 10384-92	3.2	246
1335	Carbonic anhydrase inhibitors and activators for novel therapeutic applications. *Future Medicinal Chemistry*, **2011**, 3, 1165-80	4.1	240
1334	Carbonic anhydrase inhibitors: E7070, a sulfonamide anticancer agent, potently inhibits cytosolic isozymes I and II, and transmembrane, tumor-associated isozyme IX. *Bioorganic and Medicinal Chemistry Letters*, **2004**, 14, 217-23	2.9	235
1333	Antiglaucoma carbonic anhydrase inhibitors: a patent review. *Expert Opinion on Therapeutic Patents* , **2013**, 23, 705-16	6.8	232
1332	Applications of carbonic anhydrase inhibitors and activators in therapy. *Expert Opinion on Therapeutic Patents*, **2002**, 12, 217-242	6.8	228
1331	Biochemical characterization of CA IX, one of the most active carbonic anhydrase isozymes. *Journal of Biological Chemistry*, **2008**, 283, 27799-27809	5.4	224
1330	Diuretics with carbonic anhydrase inhibitory action: a patent and literature review (2005 - 2013). *Expert Opinion on Therapeutic Patents*, **2013**, 23, 681-91	6.8	218
1329	The role of carbonic anhydrase 9 in regulating extracellular and intracellular ph in three-dimensional tumor cell growths. *Journal of Biological Chemistry*, **2009**, 284, 20299-310	5.4	218

1328	Exploiting the hydrophobic and hydrophilic binding sites for designing carbonic anhydrase inhibitors. *Expert Opinion on Drug Discovery*, **2013**, 8, 793-810	6.2	215
1327	Antiobesity carbonic anhydrase inhibitors: a literature and patent review. *Expert Opinion on Therapeutic Patents*, **2013**, 23, 725-35	6.8	213
1326	Sulfa and trimethoprim-like drugs - antimetabolites acting as carbonic anhydrase, dihydropteroate synthase and dihydrofolate reductase inhibitors. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2014**, 29, 379-87	5.6	212
1325	A small-molecule drug conjugate for the treatment of carbonic anhydrase IX expressing tumors. *Angewandte Chemie - International Edition*, **2014**, 53, 4231-5	16.4	210
1324	Carbonic anhydrase inhibitors: SAR and X-ray crystallographic study for the interaction of sugar sulfamates/sulfamides with isozymes I, II and IV. *Bioorganic and Medicinal Chemistry Letters*, **2003**, 13, 841-5	2.9	209
1323	Anticancer carbonic anhydrase inhibitors: a patent review (2008 - 2013). *Expert Opinion on Therapeutic Patents*, **2013**, 23, 737-49	6.8	208
1322	Glycosyl coumarin carbonic anhydrase IX and XII inhibitors strongly attenuate the growth of primary breast tumors. *Journal of Medicinal Chemistry*, **2011**, 54, 8271-7	8.3	201
1321	Bacterial carbonic anhydrases as drug targets: toward novel antibiotics?. *Frontiers in Pharmacology*, **2011**, 2, 34	5.6	201
1320	Carbonic anhydrase inhibitors. Inhibition of the transmembrane isozyme XII with sulfonamides-a new target for the design of antitumor and antiglaucoma drugs?. *Bioorganic and Medicinal Chemistry Letters*, **2005**, 15, 963-9	2.9	199
1319	Highly active antiretroviral therapy: current state of the art, new agents and their pharmacological interactions useful for improving therapeutic outcome. *Current Pharmaceutical Design*, **2005**, 11, 1805-43	3.3	199
1318	Dithiocarbamates strongly inhibit carbonic anhydrases and show antiglaucoma action in vivo. *Journal of Medicinal Chemistry*, **2012**, 55, 1721-30	8.3	195
1317	Carbonic anhydrase inhibitors. Inhibition of human erythrocyte isozymes I and II with a series of antioxidant phenols. *Bioorganic and Medicinal Chemistry*, **2009**, 17, 3207-11	3.4	194
1316	Diuretics: from classical carbonic anhydrase inhibitors to novel applications of the sulfonamides. *Current Pharmaceutical Design*, **2008**, 14, 641-8	3.3	194
1315	Anti-infective carbonic anhydrase inhibitors: a patent and literature review. *Expert Opinion on Therapeutic Patents*, **2013**, 23, 693-704	6.8	192
1314	Carbonic anhydrase in the scleractinian coral Stylophora pistillata: characterization, localization, and role in biomineralization. *Journal of Biological Chemistry*, **2008**, 283, 25475-25484	5.4	192
1313	Carbonic anhydrase inhibitors. Design of fluorescent sulfonamides as probes of tumor-associated carbonic anhydrase IX that inhibit isozyme IX-mediated acidification of hypoxic tumors. *Journal of Medicinal Chemistry*, **2005**, 48, 4834-41	8.3	192
1312	Carbonic anhydrase inhibitors. Inhibition of mammalian isoforms I-XIV with a series of natural product polyphenols and phenolic acids. *Bioorganic and Medicinal Chemistry*, **2010**, 18, 2159-2164	3.4	190
1311	Characterization of CA XIII, a novel member of the carbonic anhydrase isozyme family. *Journal of Biological Chemistry*, **2004**, 279, 2719-27	5.4	187

1310	Carbonic anhydrase inhibitors: X-ray and molecular modeling study for the interaction of a fluorescent antitumor sulfonamide with isozyme II and IX. *Journal of the American Chemical Society*, **2006**, 128, 8329-35	16.4	186
1309	Polyamines inhibit carbonic anhydrases by anchoring to the zinc-coordinated water molecule. *Journal of Medicinal Chemistry*, **2010**, 53, 5511-22	8.3	184
1308	Selective hydrophobic pocket binding observed within the carbonic anhydrase II active site accommodate different 4-substituted-ureido-benzenesulfonamides and correlate to inhibitor potency. *Chemical Communications*, **2010**, 46, 8371-3	5.8	180
1307	The Warburg Effect and the Hallmarks of Cancer. *Anti-Cancer Agents in Medicinal Chemistry*, **2017**, 17, 164-170	2.2	179
1306	Carbonic anhydrase inhibitors as anticonvulsant agents. *Current Topics in Medicinal Chemistry*, **2007**, 7, 855-64	3	178
1305	Structure and inhibition of the CO2-sensing carbonic anhydrase Can2 from the pathogenic fungus Cryptococcus neoformans. *Journal of Molecular Biology*, **2009**, 385, 1207-20	6.5	176
1304	The alpha and beta classes carbonic anhydrases from Helicobacter pylori as novel drug targets. *Current Pharmaceutical Design*, **2008**, 14, 622-30	3.3	175
1303	Sulfocoumarins (1,2-benzoxathiine-2,2-dioxides): a class of potent and isoform-selective inhibitors of tumor-associated carbonic anhydrases. *Journal of Medicinal Chemistry*, **2013**, 56, 293-300	8.3	174
1302	(In)organic anions as carbonic anhydrase inhibitors. *Journal of Inorganic Biochemistry*, **2012**, 111, 117-29	4.2	173
1301	Imaging of CA IX with fluorescent labelled sulfonamides distinguishes hypoxic and (re)-oxygenated cells in a xenograft tumour model. *Radiotherapy and Oncology*, **2009**, 92, 423-8	5.3	173
1300	Tumor-associated carbonic anhydrase 9 spatially coordinates intracellular pH in three-dimensional multicellular growths. *Journal of Biological Chemistry*, **2008**, 283, 20473-83	5.4	172
1299	Carbonic anhydrases as drug targets--an overview. *Current Topics in Medicinal Chemistry*, **2007**, 7, 825-33	3	171
1298	Carbonic anhydrase inhibitors: interactions of phenols with the 12 catalytically active mammalian isoforms (CA I-XIV). *Bioorganic and Medicinal Chemistry Letters*, **2008**, 18, 1583-7	2.9	170
1297	Carbonic anhydrase inhibitors: inhibition of isozymes I, II, and IX with triazole-linked O-glycosides of benzene sulfonamides. *Journal of Medicinal Chemistry*, **2007**, 50, 1651-7	8.3	169
1296	Sulfamates and their therapeutic potential. *Medicinal Research Reviews*, **2005**, 25, 186-228	14.4	169
1295	Sulfonamides: a patent review (2008 - 2012). *Expert Opinion on Therapeutic Patents*, **2012**, 22, 747-58	6.8	167
1294	Carbonic anhydrase inhibitors. Zonisamide is an effective inhibitor of the cytosolic isozyme II and mitochondrial isozyme V: solution and X-ray crystallographic studies. *Bioorganic and Medicinal Chemistry Letters*, **2005**, 15, 2315-20	2.9	166
1293	Synthesis and carbonic anhydrase isoenzymes I, II, IX, and XII inhibitory effects of dimethoxybromophenol derivatives incorporating cyclopropane moieties. *Journal of Medicinal Chemistry*, **2015**, 58, 640-50	8.3	164

1292	Carbonic anhydrase inhibitors. The mitochondrial isozyme VB as a new target for sulfonamide and sulfamate inhibitors. *Journal of Medicinal Chemistry*, **2005**, 48, 7860-6	8.3	161
1291	Inhibition of carbonic anhydrase IX targets primary tumors, metastases, and cancer stem cells: Three for the price of one. *Medicinal Research Reviews*, **2018**, 38, 1799-1836	14.4	159
1290	In vitro inhibition of □carbonic anhydrase isozymes by some phenolic compounds. *Bioorganic and Medicinal Chemistry Letters*, **2011**, 21, 4259-62	2.9	158
1289	In Vitro inhibition of human carbonic anhydrase I and II isozymes with natural phenolic compounds. *Chemical Biology and Drug Design*, **2011**, 77, 494-9	2.9	154
1288	Bacterial, fungal and protozoan carbonic anhydrases as drug targets. *Expert Opinion on Therapeutic Targets*, **2015**, 19, 1689-704	6.4	153
1287	Taking advantage of tumor cell adaptations to hypoxia for developing new tumor markers and treatment strategies. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2009**, 24 Suppl 1, 1-39	5.6	153
1286	Therapeutic potential of sulfamides as enzyme inhibitors. *Medicinal Research Reviews*, **2006**, 26, 767-92	14.4	153
1285	A novel class of carbonic anhydrase inhibitors: glycoconjugate benzene sulfonamides prepared by "click-tailing". *Journal of Medicinal Chemistry*, **2006**, 49, 6539-48	8.3	153
1284	Carbonic Anhydrase Inhibition and the Management of Hypoxic Tumors. *Metabolites*, **2017**, 7,	5.6	151
1283	Carbonic anhydrase IX: Biochemical and crystallographic characterization of a novel antitumor target. *Biochimica Et Biophysica Acta - Proteins and Proteomics*, **2010**, 1804, 404-9	4	151
1282	Carbonic anhydrase inhibitors: stacking with Phe131 determines active site binding region of inhibitors as exemplified by the X-ray crystal structure of a membrane-impermeant antitumor sulfonamide complexed with isozyme II. *Journal of Medicinal Chemistry*, **2005**, 48, 5721-7	8.3	150
1281	Dithiocarbamates: a new class of carbonic anhydrase inhibitors. Crystallographic and kinetic investigations. *Chemical Communications*, **2012**, 48, 1868-70	5.8	149
1280	Modulation of carbonic anhydrase activity and its applications in therapy. *Expert Opinion on Therapeutic Patents*, **2004**, 14, 667-702	6.8	148
1279	Carbonic anhydrase inhibitors: synthesis of water-soluble, topically effective intraocular pressure lowering aromatic/heterocyclic sulfonamides containing 8-quinoline-sulfonyl moieties: is the tail more important than the ring?. *Bioorganic and Medicinal Chemistry*, **1999**, 7, 2397-406	3.4	148
1278	Carbonic anhydrase inhibitors: the beta-carbonic anhydrase from Helicobacter pylori is a new target for sulfonamide and sulfamate inhibitors. *Bioorganic and Medicinal Chemistry Letters*, **2007**, 17, 3585-94	2.9	146
1277	Indisulam: an anticancer sulfonamide in clinical development. *Expert Opinion on Investigational Drugs*, **2003**, 12, 283-7	5.9	146
1276	Carbonic anhydrase inhibitors: synthesis of water-soluble, aminoacyl/dipeptidyl sulfonamides possessing long-lasting intraocular pressure-lowering properties via the topical route. *Journal of Medicinal Chemistry*, **1999**, 42, 3690-700	8.3	146
1275	Saccharin inhibits carbonic anhydrases: possible explanation for its unpleasant metallic aftertaste. *Angewandte Chemie - International Edition*, **2007**, 46, 7697-9	16.4	145

1274	Carbonic anhydrase inhibitors. Design of selective, membrane-impermeant inhibitors targeting the human tumor-associated isozyme IX. *Journal of Medicinal Chemistry*, **2004**, 47, 2337-47	8.3	145
1273	Specific inhibition of carbonic anhydrase IX activity enhances the in vivo therapeutic effect of tumor irradiation. *Radiotherapy and Oncology*, **2011**, 99, 424-31	5.3	144
1272	Nonaromatic sulfonamide group as an ideal anchor for potent human carbonic anhydrase inhibitors: role of hydrogen-bonding networks in ligand binding and drug design. *Journal of Medicinal Chemistry*, **2002**, 45, 3583-7	8.3	144
1271	7,8-disubstituted- but not 6,7-disubstituted coumarins selectively inhibit the transmembrane, tumor-associated carbonic anhydrase isoforms IX and XII over the cytosolic ones I and II in the low nanomolar/subnanomolar range. *Bioorganic and Medicinal Chemistry Letters*, **2010**, 20, 7255-8	2.9	143
1270	Carbonic anhydrase inhibitors and activators and their use in therapy. *Expert Opinion on Therapeutic Patents*, **2006**, 16, 1627-1664	6.8	143
1269	New strategies for targeting the hypoxic tumour microenvironment in breast cancer. *Cancer Treatment Reviews*, **2013**, 39, 171-9	14.4	142
1268	In vitro inhibition of salicylic acid derivatives on human cytosolic carbonic anhydrase isozymes I and II. *Bioorganic and Medicinal Chemistry*, **2008**, 16, 9101-5	3.4	142
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1266	Direct extracellular interaction between carbonic anhydrase IV and the human NBC1 sodium/bicarbonate co-transporter. *Biochemistry*, **2003**, 42, 12321-9	3.2	142
1265	Carbonic anhydrase inhibitors: the first selective, membrane-impermeant inhibitors targeting the tumor-associated isozyme IX. *Bioorganic and Medicinal Chemistry Letters*, **2004**, 14, 869-73	2.9	140
1264	Carbonic anhydrase inhibitors - Part 49: Synthesis of substituted ureido and thioureido derivatives of aromatic/heterocyclic sulfonamides with increased affinities for isozyme I. *European Journal of Medicinal Chemistry*, **1998**, 33, 83-93	6.8	139
1263	Carbonic anhydrase inhibitors as emerging drugs for the treatment of obesity. *Expert Opinion on Emerging Drugs*, **2008**, 13, 383-92	3.7	139
1262	Carbonic anhydrase inhibitors as emerging agents for the treatment and imaging of hypoxic tumors. *Expert Opinion on Investigational Drugs*, **2018**, 27, 963-970	5.9	139
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1260	Imaging the hypoxia surrogate marker CA IX requires expression and catalytic activity for binding fluorescent sulfonamide inhibitors. *Radiotherapy and Oncology*, **2007**, 83, 367-73	5.3	138
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1257	Carbonic anhydrase inhibitors: DNA cloning and inhibition studies of the alpha-carbonic anhydrase from Helicobacter pylori, a new target for developing sulfonamide and sulfamate gastric drugs. *Journal of Medicinal Chemistry*, **2006**, 49, 2117-26	8.3	137

1256	Carbonic anhydrase inhibitors: X-ray crystallographic structure of the adduct of human isozyme II with EMATE, a dual inhibitor of carbonic anhydrases and steroid sulfatase. *Bioorganic and Medicinal Chemistry Letters*, **2004**, 14, 231-4	2.9	137
1255	Carbonic anhydrase inhibitors: water-soluble 4-sulfamoylphenylthioureas as topical intraocular pressure-lowering agents with long-lasting effects. *Journal of Medicinal Chemistry*, **2000**, 43, 4884-92	8.3	137
1254	The class carbonic anhydrases as drug targets for antimalarial agents. *Expert Opinion on Therapeutic Targets*, **2015**, 19, 551-63	6.4	135
1253	Carbonic anhydrase inhibitors. Antioxidant polyphenols effectively inhibit mammalian isoforms I-XV. *Bioorganic and Medicinal Chemistry Letters*, **2010**, 20, 5050-3	2.9	135
1252	Carbonic anhydrase inhibitors in the treatment and prophylaxis of obesity. *Expert Opinion on Therapeutic Patents*, **2003**, 13, 1545-1550	6.8	135
1251	Rosmarinic acid inhibits some metabolic enzymes including glutathione S-transferase, lactoperoxidase, acetylcholinesterase, butyrylcholinesterase and carbonic anhydrase isoenzymes. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2016**, 31, 1698-702	5.6	134
1250	Sulfonamides and their isosters as carbonic anhydrase inhibitors. *Future Medicinal Chemistry*, **2014**, 6, 1149-65	4.1	133
1249	Carbonic anhydrase inhibitors. Inhibition of tumor-associated isozyme IX by halogenosulfanilamide and halogenophenylaminobenzolamide derivatives. *Journal of Medicinal Chemistry*, **2003**, 46, 2187-96	8.3	133
1248	Carbonic anhydrase inhibitors Part 29 1: Interaction of isozymes I, II and IV with benzolamide-like derivatives. *European Journal of Medicinal Chemistry*, **1998**, 33, 739-751	6.8	130
1247	Metal binding and antibacterial activity of ciprofloxacin complexes. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2005**, 20, 303-7	5.6	130
1246	Zinc complexes of benzothiazole-derived Schiff bases with antibacterial activity. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2003**, 18, 259-63	5.6	130
1245	Carbonic anhydrase inhibitors: anticonvulsant sulfonamides incorporating valproyl and other lipophilic moieties. *Journal of Medicinal Chemistry*, **2002**, 45, 312-20	8.3	129
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1241	Carbonic anhydrase inhibitors. Inhibition of the human cytosolic isozyme VII with aromatic and heterocyclic sulfonamides. *Bioorganic and Medicinal Chemistry Letters*, **2005**, 15, 971-6	2.9	128
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1237	Metal-Based Antibacterial and Antifungal Agents: Synthesis, Characterization, and In Vitro Biological Evaluation of Co(II), Cu(II), Ni(II), and Zn(II) Complexes With Amino Acid-Derived Compounds. *Bioinorganic Chemistry and Applications*, **2006**, 2006, 83131	4.2	124
1236	Antimetastatic effect of sulfamate carbonic anhydrase IX inhibitors in breast carcinoma xenografts. *Journal of Medicinal Chemistry*, **2012**, 55, 5591-600	8.3	123
1235	Carbonic anhydrase activators. Activation of isozymes I, II, IV, VA, VII, and XIV with l- and d-histidine and crystallographic analysis of their adducts with isoform II: engineering proton-transfer processes within the active site of an enzyme. *Chemistry - A European Journal*, **2006**, 12, 7057-66	4.8	122
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1231	Dithiocarbamates strongly inhibit the class carbonic anhydrases from Mycobacterium tuberculosis. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2013**, 28, 407-11	5.6	118
1230	Coumarins incorporating hydroxy- and chloro-moieties selectively inhibit the transmembrane, tumor-associated carbonic anhydrase isoforms IX and XII over the cytosolic ones I and II. *Bioorganic and Medicinal Chemistry Letters*, **2010**, 20, 4511-4	2.9	118
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1228	Selection of Carbonic Anhydrase IX Inhibitors from One Million DNA-Encoded Compounds. *ACS Chemical Biology*, **2011**, 6, 336-44	4.9	117
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1226	Carbonic anhydrase activators. Activation of isoforms I, II, IV, VA, VII, and XIV with L- and D-phenylalanine and crystallographic analysis of their adducts with isozyme II: stereospecific recognition within the active site of an enzyme and its consequences for the drug design. *Journal of Medicinal Chemistry*, **2006**, 49, 3019-27	8.3	116
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1224	Inhibitors of HIV-1 protease: current state of the art 10 years after their introduction. From antiretroviral drugs to antifungal, antibacterial and antitumor agents based on aspartic protease inhibitors. *Current Medicinal Chemistry*, **2007**, 14, 2734-48	4.3	115
1223	Unsymmetrical 1,1'-disubstituted ferrocenes: synthesis of Co(ii), Cu(ii), Ni(ii) and Zn(ii) chelates of ferrocenyl -1-thiadiazolo-1'-tetrazole, -1-thiadiazolo-1'-triazole and -1-tetrazolo-1'-triazole with antimicrobial properties. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2002**, 17, 261-6	5.6	115
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1220	Carbonic anhydrase inhibitors: inhibition of mammalian isoforms I-XIV with a series of substituted phenols including paracetamol and salicylic acid. *Bioorganic and Medicinal Chemistry*, **2008**, 16, 7424-8	3.4	114
1219	Inhibition of the archaeal beta-class (Cab) and gamma-class (Cam) carbonic anhydrases. *Current Topics in Medicinal Chemistry*, **2007**, 7, 901-8	3	113
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